16-1150. Applicants respectfully request that the amendments and remarks made herein be entered into the record of the instant application.

IN THE CLAIMS:

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Please amend the claims as set forth below. A marked-up copy of the claim amendments is provided as Exhibit A, and a copy of the pending claims is provided as Exhibit B.

Please amend claim 34 to read as follows:

34. (Amended) A method for treating cerebral ischemia in a human subject comprising peripherally administering to said human subject a non-toxic amount of erythropoietin effective to exert a neuroprotective effect.

Please add new claims 40 to 63 as follows:

- 40. (New) A method for treating cerebral ischemia in a mammal comprising peripherally administering to said mammal an amount of erythropoietin effective to exert a neuroprotective effect without a toxic increase in hemoglobin concentration or hematocrit.
- 41. (New) The method of Claim 40 wherein said administering is carried out in a vascular fashion.
- 42. (New) The method of Claim 40 wherein said vascular administration is intravenous.
- 43. (New) The method of Claim 40, 41, or 42 wherein said erythropoietin is administered for the treatment of stroke.
- 44. (New) The method of Claim 40, 41, or 42 wherein said erythropoietin is administered at a dosage of 50,000 to 100,000 Units per administration or per day.

- 45. (New) The method of Claim 40, 41, or 42 wherein said erythropoietin is native erythropoietin, recombinant human erythropoietin or animal erythropoietin or a derivative thereof.
- 46. (New) A method for treating cerebral ischemia in a human subject comprising peripherally administering to said human subject an amount of erythropoietin effective to exert a neuroprotective effect without a toxic increase in hemoglobin concentration or hematocrit.
- 47. (New) The method of Claim 46 wherein said administering is carried out in a vascular fashion.
- 48. (New) The method of Claim 47 wherein said vascular administration is intravenous.
- 49. (New) The method of Claim 46, 47, or 48 wherein said erythropoietin is administered for the treatment of stroke.
- 50. (New) The method of Claim 46, 47, or 48 wherein said erythropoietin is administered at a dosage of 50,000 to 100,000 Units per administration or per day.
- 51. (New) The method of Claim 46, 47, or 48 wherein said erythropoietin is native erythropoietin, recombinant human erythropoietin or animal erythropoietin or a derivative thereof.
- 52. (New) A method for treating cerebral ischemia in a mammal comprising peripherally administering to said mammal an amount of erythropoietin effective to exert a neuroprotective effect without an increase in hematocrit in said mammal.
- 53. (New) The method of Claim 52 wherein said administering is carried out in a vascular fashion.

- 54. (New) The method of Claim 53 wherein said vascular administration is intravenous.
- 55. (New) The method of Claim 52, 53, or 54 wherein said erythropoietin is administered for the treatment of stroke.
- 56. (New) The method of Claim 52, 53, or 54 wherein said erythropoietin is administered at a dosage of 50,000 to 100,000 Units per administration or per day.
- 57. (New) The method of Claim 52, 53, or 54 wherein said erythropoietin is native erythropoietin, recombinant human erythropoietin or animal erythropoietin or a derivative thereof.
- 58. (New) A method for treating cerebral ischemia in a human subject comprising peripherally administering to said human subject an amount of erythropoietin effective to exert a neuroprotective effect without an increase in hematocrit in said human subject.
- 59. (New) The method of Claim 58 wherein said administering is carried out in a vascular fashion.
- 60. (New) The method of Claim 58 wherein said vascular administration is intravenous.
- 61. (New) The method of Claim 58, 59, or 60 wherein said erythropoietin is administered for the treatment of stroke.
- 62. (New) The method of Claim 58, 59, or 60 wherein said erythropoietin is administered at a dosage of 50,000 to 100,000 Units per administration or per day.